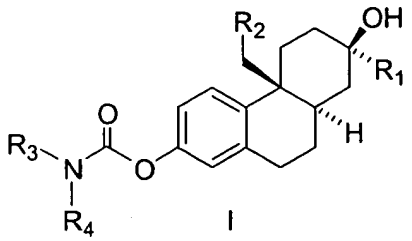
In the Claims

Please amend the claims as follows:

Claim 1 (Previously Amended) A compound of Formula I



I

or a pharmaceutically acceptable salt of said compound;

wherein R_1 is a) -(C_1-C_6)alkyl optionally substituted with -CF_3, b) -C≡C-CH_3, c) -C≡C-Cl, d) -C≡C-CF_3, e) -CH_2O(C_1-C_4)alkyl optionally substituted with -CF_3 or f) -CF_3;

R_2 is a) -(C_1-C_5)alkyl, b) -(C_2-C_5)alkenyl or c) -phenyl optionally substituted with one of the following: -OH, -NR_9-C(O)-(C_2-C_4)alkyl, -CN, -Z-het, -O-(C_1-C_3)alkyl-C(O)-NR_9R_{10}, -NR_9-Z-C(O)-NR_9R_{10}, -Z-NR_9-SO_2-R_{10}, -NR_9-SO_2-het, -O-C(O)-(C_1-C_4)alkyl or -O-SO_2-(C_1-C_4)alkyl;

Z for each occurrence is independently -(C_0-C_4)alkyl;

R_3 is a) -hydrogen, b) -(C_1-C_6)alkyl optionally substituted with one to three halo, c) -(C_2-C_6)alkenyl or d) -(C_2-C_6)alkynyl optionally substituted with one to three halo;

R_4 is a) -hydrogen, or b) -(C_2-C_5)alkyl-NR_5R_6;

R_5 and R_6 are each independently a) hydrogen or b) -(C_1-C_3)alkyl;

het is an optionally substituted 5-, 6- or 7-membered saturated, partially saturated or unsaturated heterocyclic ring containing from 1 to 3 heteroatoms selected from the group consisting of nitrogen, oxygen and sulfur; and including any bicyclic group in which any of the above heterocyclic rings is fused to a benzene ring or another heterocyclic ring; and optionally

substituted with one to four R_7; provided that het is other than pyridinyl, imidazolyl or tetrazolyl;

R_7 is a) -(C_1-C_6)alkyl optionally substituted with one to three R_8, b) –Z-NR_9R_{10} or c) –Z-C(O)-NR_9R_{10};

R_8 for each occurrence is independently a) halo, b) –OH, c) oxo or d) -O(C_1-C_6)alkyl;

R_9 and R_{10} for each occurrence are independently a) -H or b) -(C_1-C_3)alkyl;

or R_9 and R_{10} are taken together with N to form het;

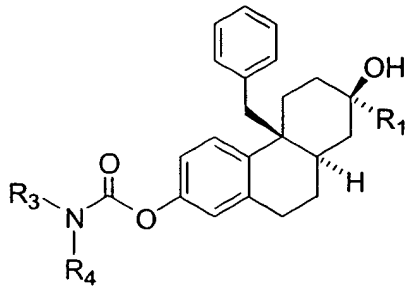
provided that:

1) when R_1 is –C≡C-CH_3, R_2 is phenyl and R_3 is hydrogen, then R_4 is other than –$(CH_2)_2$-$N(CH_3)_2$, or –$(CH_2)_3$-$N(CH_3)_2$, –$(CH_2)_2$-pyrrolidinyl optionally substituted with methyl, –$(CH_2)_3$-pyrrolidinyl or –$(CH_2)_2$-morpholinyl;

2) when R_1 is –C≡C-CH_3, R_2 is propyl and R_3 is hydrogen, then R_4 is other than –$(CH_2)_2$-$N(CH_3)_2$; and

3) when R_1 is –C≡C-CH_3, R_2 is butyl and R_3 is hydrogen, then R_4 is other than –$(CH_2)_2$-$N(CH_3)_2$.

Claim 2 (Previously Amended) A compound of claim 1 of Formula II



II

or a pharmaceutically acceptable salt of said compound;

wherein R_1 is a) -(C_1-C_6)alkyl optionally substituted with -CF_3, b) -C≡C-CH_3, c) –CF_3 or d) –$CH_2O(C_2$-C_4)alkyl.

Claim 3 (Original) A compound of claim 2 wherein R_1 is a) –$CH_2CH_2CH_3$, b) -C≡C-CH_3 or c) –CF_3.

Claim 4 (Original) A compound of claim 3

wherein R_3 is a) hydrogen, b) methyl, c) ethyl, d) propyl or e) isopropyl;

R_4 is -(C_2-C_3)alkyl-NR_5R_6;

R_5 and R_6 are each independently a) methyl, b) ethyl, c) propyl or d) isopropyl.

Claim 5 (Original) A compound of claim 4

wherein R_3 is a) methyl, b) ethyl, c) propyl or d) isopropyl;

R_4 is -(C_2-C_3)alkyl-NR_5R_6;

R_5 and R_6 are each independently a) methyl, b) ethyl, c) propyl or d) isopropyl.

Claim 6 (Original) A compound of claim 5

wherein R_3 is a) methyl or b) ethyl;

R_4 is -(C_2-C_3)alkyl-NR_5R_6;

R_5 and R_6 are each methyl.

Claims 7-11 (Previously Canceled)

Claim 12 (Original) A compound of claim 1

wherein R_1 is a) $-CH_2CH_2CH_3$, b) $-C\equiv C-CH_3$ or c) $-CF_3$;

R_2 is a) -(C_1-C_5)alkyl or b) -(C_2-C_5)alkenyl;

R_3 is a) hydrogen, b) methyl, c) ethyl, d) propyl or e) isopropyl;

R_4 is -(C_2-C_3)alkyl-NR_5R_6;

R_5 and R_6 are each independently a) methyl, b) ethyl, c) propyl or d) isopropyl.

Claim13 (Original) A compound of claim 12

wherein R_2 is a) methyl, b) ethyl, c) propyl, d) ethenyl, e) propenyl or f) butenyl;

R_3 is a) hydrogen, b) methyl or c) ethyl,

R_5 and R_6 are each independently a) methyl or b) ethyl.

Claims 14-17 (Previously Canceled)

Claim 18 (Original) A compound of claim 1 wherein in Formula I $-CH_2-R_2$ is ethenyl or ethynyl.

Claim 19 (Original) A compound of claim 4 selected from the group consisting of:

carbamic acid, [2-(dimethylamino)ethyl]-, (4b*S*,7*R*,8a*R*)-4b,5,6,7,8,8a,9,10-octahydro-7-hydroxy-4b-(phenylmethyl)-7-(trifluoromethyl)-2-phenanthrenyl ester;

carbamic acid, [3-(dimethylamino)propyl]-, (4b*S*,7*R*,8a*R*)-4b,5,6,7,8,8a,9,10-octahydro-7-hydroxy-4b-(phenylmethyl)-7-(trifluoromethyl)-2-phenanthrenyl ester; and

carbamic acid, [3-(diethylamino)propyl]-, (4b*S*,7*R*,8a*R*)-4b,5,6,7,8,8a,9,10-octahydro-7-hydroxy-4b-(phenylmethyl)-7-(trifluoromethyl)-2-phenanthrenyl ester.

Claim 20 (Original) A compound of claim 6 selected from the group consisting of:

carbamic acid, [2-(dimethylamino)ethyl]methyl-, (4b*S*,7*R*,8a*R*)-4b,5,6,7,8,8a,9,10-octahydro-7-hydroxy-4b-(phenylmethyl)-7-(trifluoromethyl)-2-phenanthrenyl ester;

carbamic acid, [2-(dimethylamino)ethyl]methyl-, (4b*S*,7*R*,8a*R*)-4b,5,6,7,8,8a,9,10-octahydro-7-hydroxy-4b-(phenylmethyl)-7-propyl-2-phenanthrenyl ester;

carbamic acid, [3-(dimethylamino)propyl]ethyl-, (4b*S*,7*R*,8a*R*)-4b,5,6,7,8,8a,9,10-octahydro-7-hydroxy-4b-(phenylmethyl)-7-(trifluoromethyl)-2-phenanthrenyl ester; and

carbamic acid, [2-(dimethylamino)ethyl]ethyl-, (4b*S*,7*R*,8a*R*)-4b,5,6,7,8,8a,9,10-octahydro-7-hydroxy-4b-(phenylmethyl)-7-(trifluoromethyl)-2-phenanthrenyl ester.

Claim 21-23 (Previously Canceled)

Claim 24 (Original) A compound of claim 13 selected from the group consisting of:

carbamic acid, (3-dimethylaminopropyl)methyl-, (4b*S*, 7*R*, 8a*R*)-4b,5,6,7,8,8a,9,10-octahydro-4b-ethyl-7-hydroxy-7-prop-1-ynyl-phenanthren-2-yl ester;

carbamic acid, (2-dimethylaminoethyl)methyl-, (4b*S*, 7*R*, 8a*R*)-4b,5,6,7,8,8a,9,10-octahydro-4b-ethyl-7-hydroxy-7-prop-1-ynyl-phenanthren-2-yl ester;

carbamic acid, (2-dimethylaminoethyl)ethyl-, (4b*S*, 7*R*, 8a*R*)-4b,5,6,7,8,8a,9,10-octahydro-4b-ethyl-7-hydroxy-7-prop-1-ynyl-phenanthren-2-yl ester; and

carbamic acid, (2-dimethylaminoethyl)-, (4b*S*, 7*R*, 8a*R*)-4b,5,6,7,8,8a,9,10-octahydro-4b-ethyl-7-hydroxy-7-prop-1-ynyl-phenanthren-2-yl ester.

Claims 25-26 (Previously canceled)

Claim 27 (Newly amended) A method for the treatment of a glucocorticoid receptor-mediated disease or condition which is selected from **the group consisting of** obesity, diabetes, depression, anxiety and neurodegeneration in a mammal, which comprises administering to the mammal a therapeutically effective amount of a compound of claim 1, or a pharmaceutically acceptable salt of said compound.

Claim 28 (Previously canceled)

Claim 29 (Previously amended) The method of claim 27 wherein the condition is obesity.

Claims 30-41 (Previously canceled)